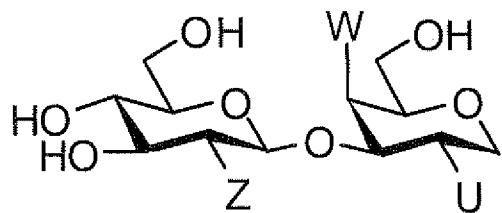


AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Claims 1-69 (Cancelled).

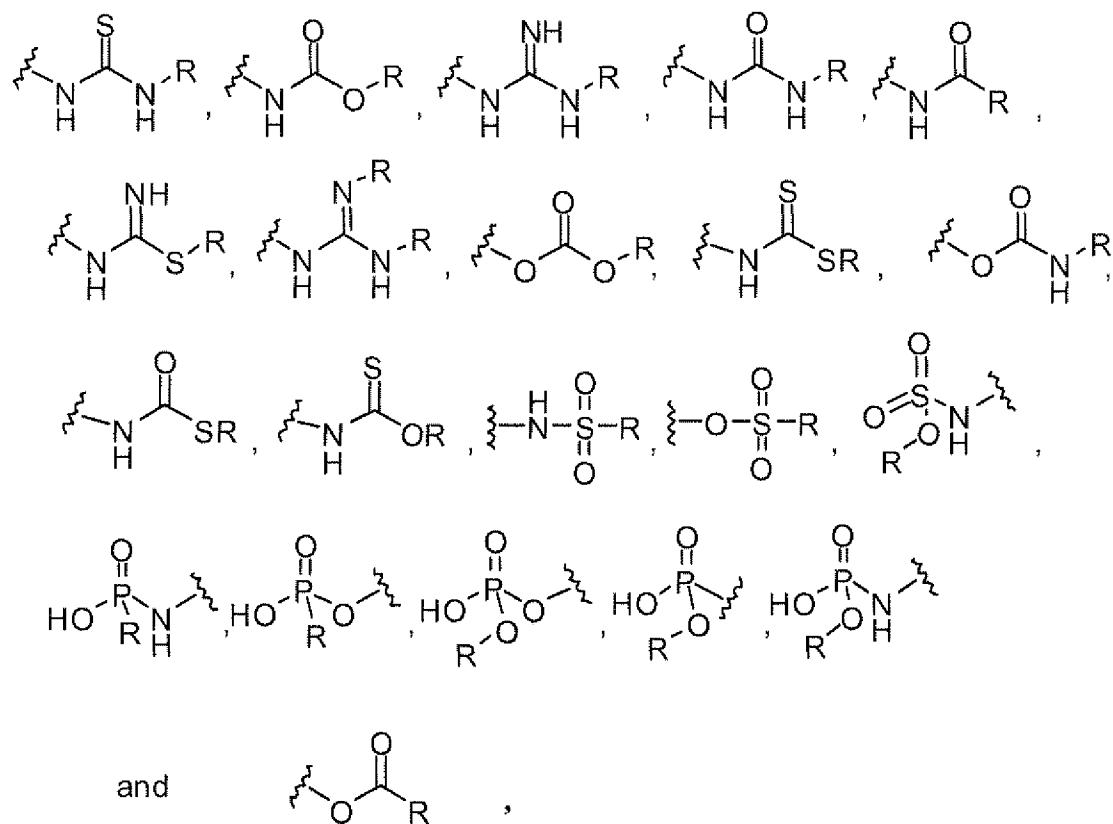
70. (New) A method of inhibiting bacterial growth comprising contacting a bacteria with at least one disaccharide compound of General Formula (I),



General Formula (I)

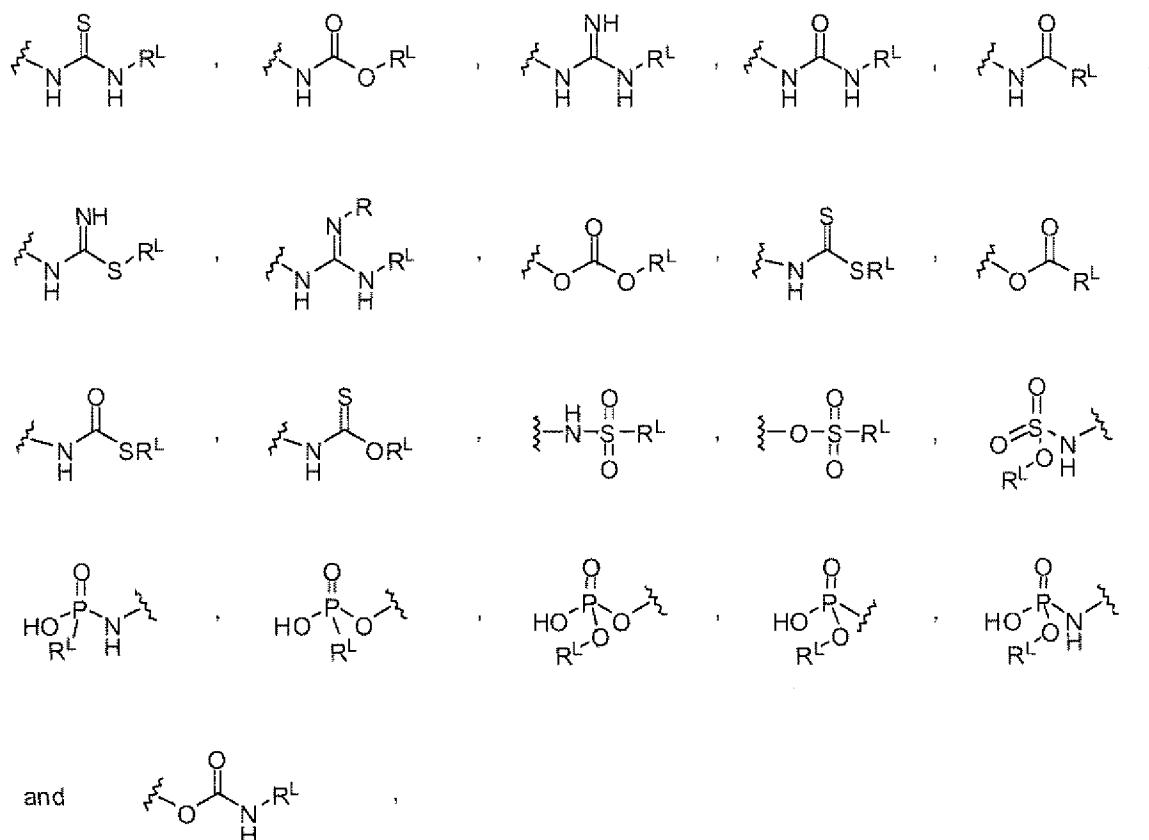
wherein

U and Z are independently selected from the group consisting of: -OR, -NHR, -NR(R),



wherein R may be the same or different, R is a moiety of not more than 20 carbon atoms independently selected from the group consisting of: alkyl, alkenyl, alkynyl, heteroalkyl, aryl, heteroaryl, arylalkyl and heteroarylalkyl;

W is independently selected from the group consisting of -OR^L, -NHR^L, -NR^LR,

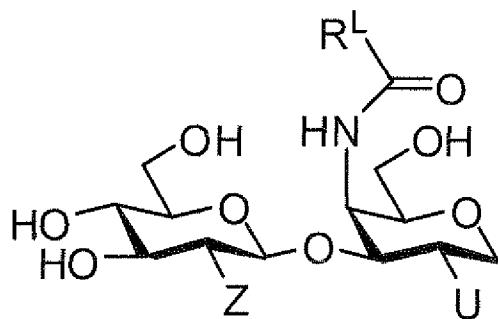


wherein R^L is a substituted or unsubstituted, linear or branched moiety of between 3 and 55 carbon atoms selected from the group consisting of: alkyl, heteroalkyl, arylalkyl, and alkylaryl chain.

71. (New) The method of claim 70, wherein R^L is substituted by a moiety selected from the group consisting of: carboxylic acids, sulfonic acids, phosphoric acids, tetrazoles, amines, guanidiniums, amidines, imidazoles, and oxazoles.

72. (New) The method of claim 70, wherein one or more R groups is substituted by a moiety selected from the group consisting of: -OH, -NO, -NO₂, -NH₂, -N₃, halogen, -CF₃, -CHF₂, -CH₂F, nitrile, alkoxy, aryloxy, amidine, guanidiniums, carboxylic acid, carboxylic acid ester, carboxylic acid amide, aryl, cycloalkyl, heteroalkyl, heteroaryl, aminoalkyl, aminodialkyl, aminotrialkyl, aminoacyl, carbonyl, substituted or unsubstituted imine, sulfate, sulfonamide, phosphate, phosphoramido, hydrazide, hydroxamate, hydroxamic acid, heteroaryloxy, carbamoyl, aminoaryl, aminoheteroaryl, thioalkyl, thioaryl and thioheteroaryl.

73. (New) The method of claim 70, wherein the compound is of General Formula (III):



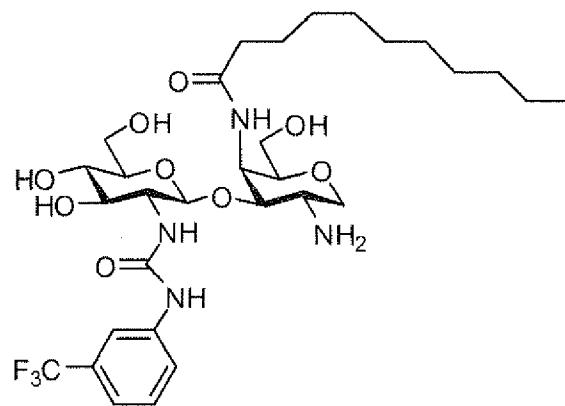
General Formula (III).

74. (New) The method of claim 70, wherein the bacteria is a Gram + bacteria.

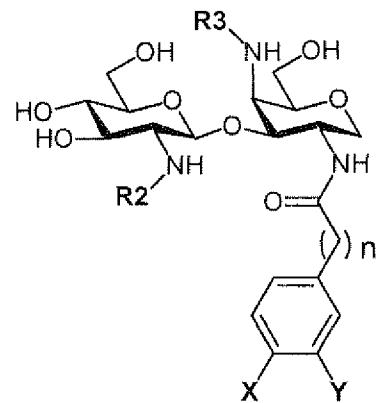
75. (New) The method of claim 70, wherein the bacteria is a Gram – bacteria.

76. (New) The method of claim 70, wherein the bacteria is selected from the group consisting of an *Escherichia coli* (*E. coli*), *Micrococcus luteus*, *Staphylococcus aureus*, Methicillin-resistant *Staphylococcus aureus* (MRSA), *Enterococcus faecalis*, *Enterococcus faecalis* Vancomycin resistant and *Streptococcus pyogenes*.

77. (New) The method of claim 70, wherein the bacteria is *Staphylococcus aureus* and the compound is

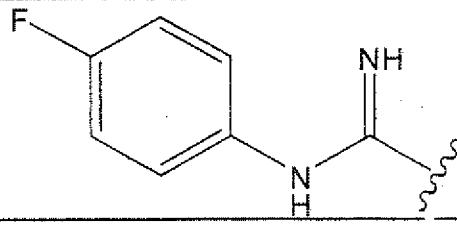
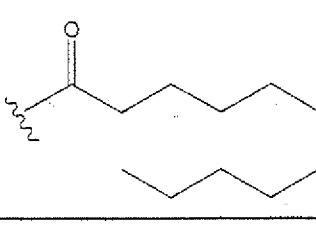
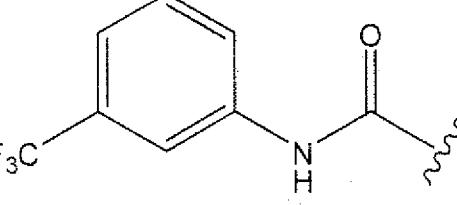
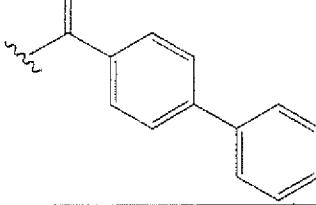
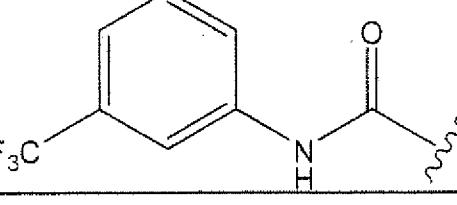
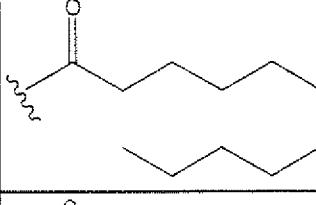
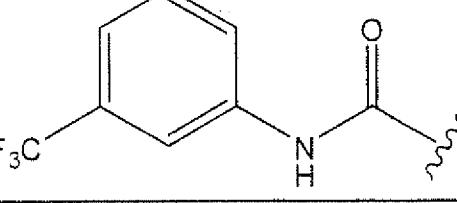
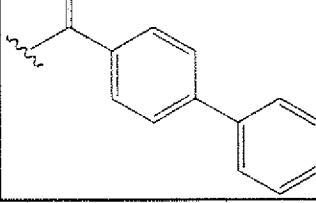


78. (New) The method of claim 70, wherein the bacteria is *Staphylococcus aureus* and the compound is

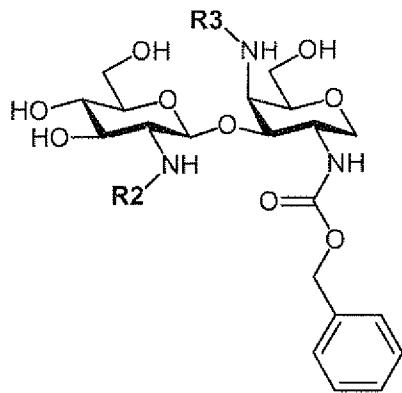


wherein:

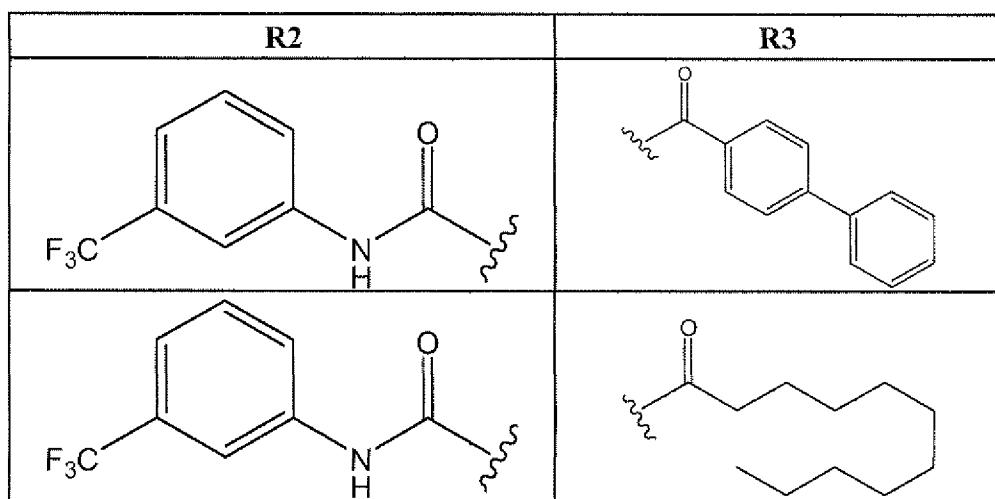
n	X	Y	R2	R3
1	H	CF ₃		
1	H	CF ₃		

0	H	CF ₃		
0	H	CF ₃		
0	H	CF ₃		
I	CF ₃	H		

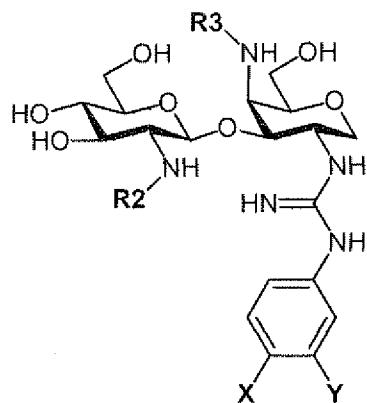
79. (New) The method of claim 70, wherein the bacteria is *Staphylococcus aureus* and the compound is



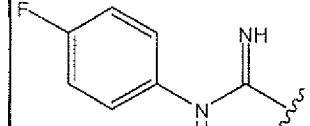
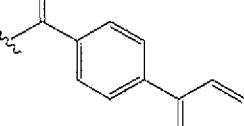
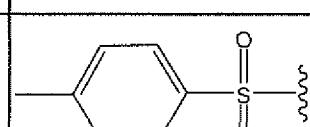
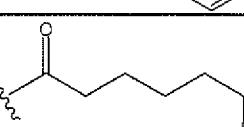
wherein:

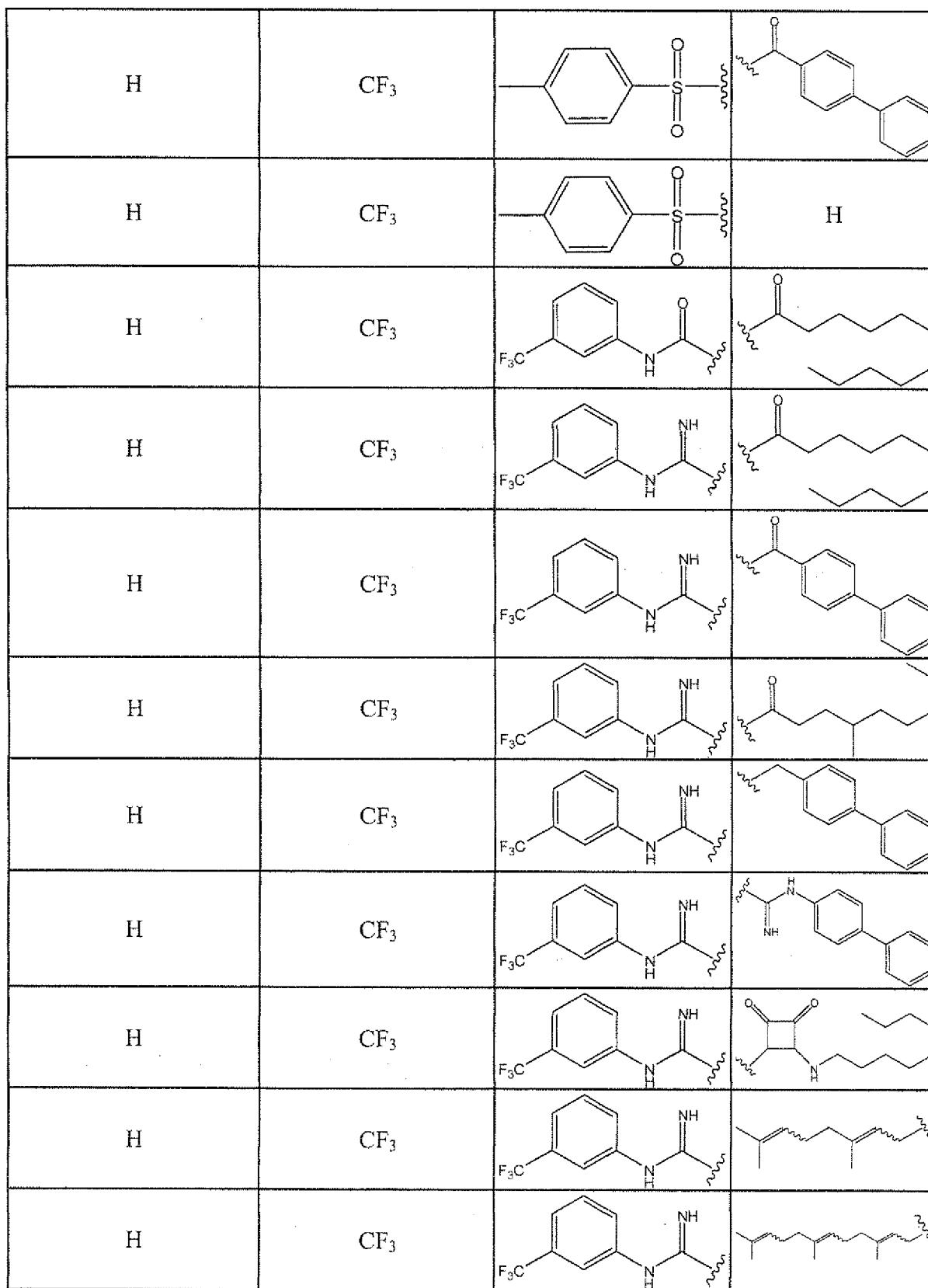


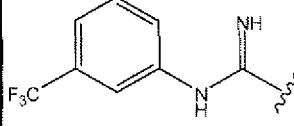
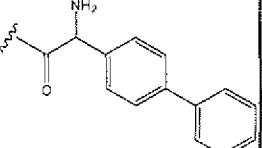
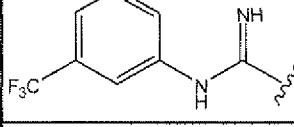
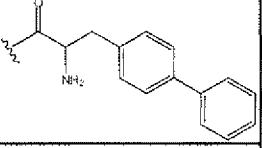
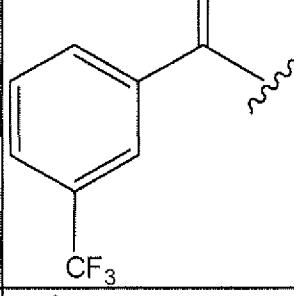
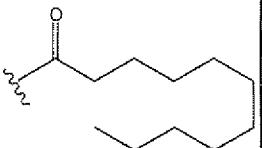
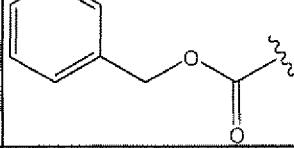
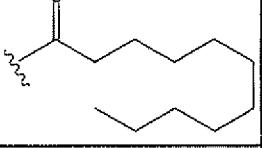
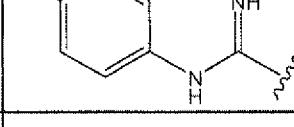
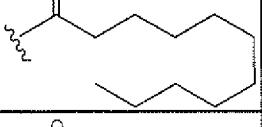
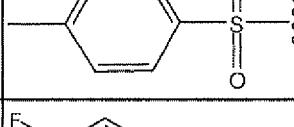
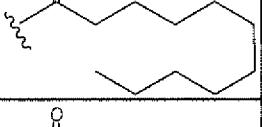
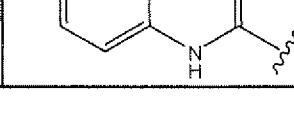
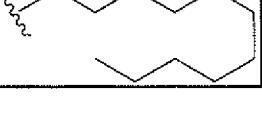
80. (New) The method of claim 70, wherein the bacteria is *Staphylococcus aureus* and the compound is



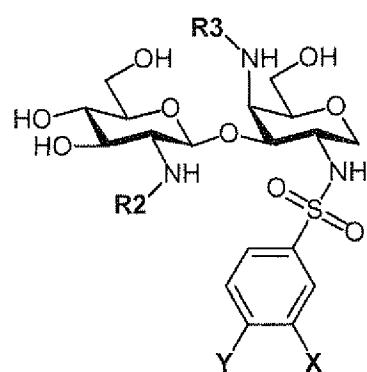
wherein:

X	Y	R2	R3
H	CF ₃		
H	CF ₃		



H	CF ₃		
H	CF ₃		
F	H		
F	H		
F	H		
F	H		
F	H		

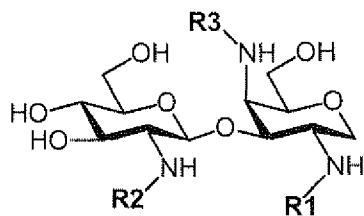
81. (New) The method of claim 70, wherein the bacteria is *Staphylococcus aureus* and the compound is



wherein:

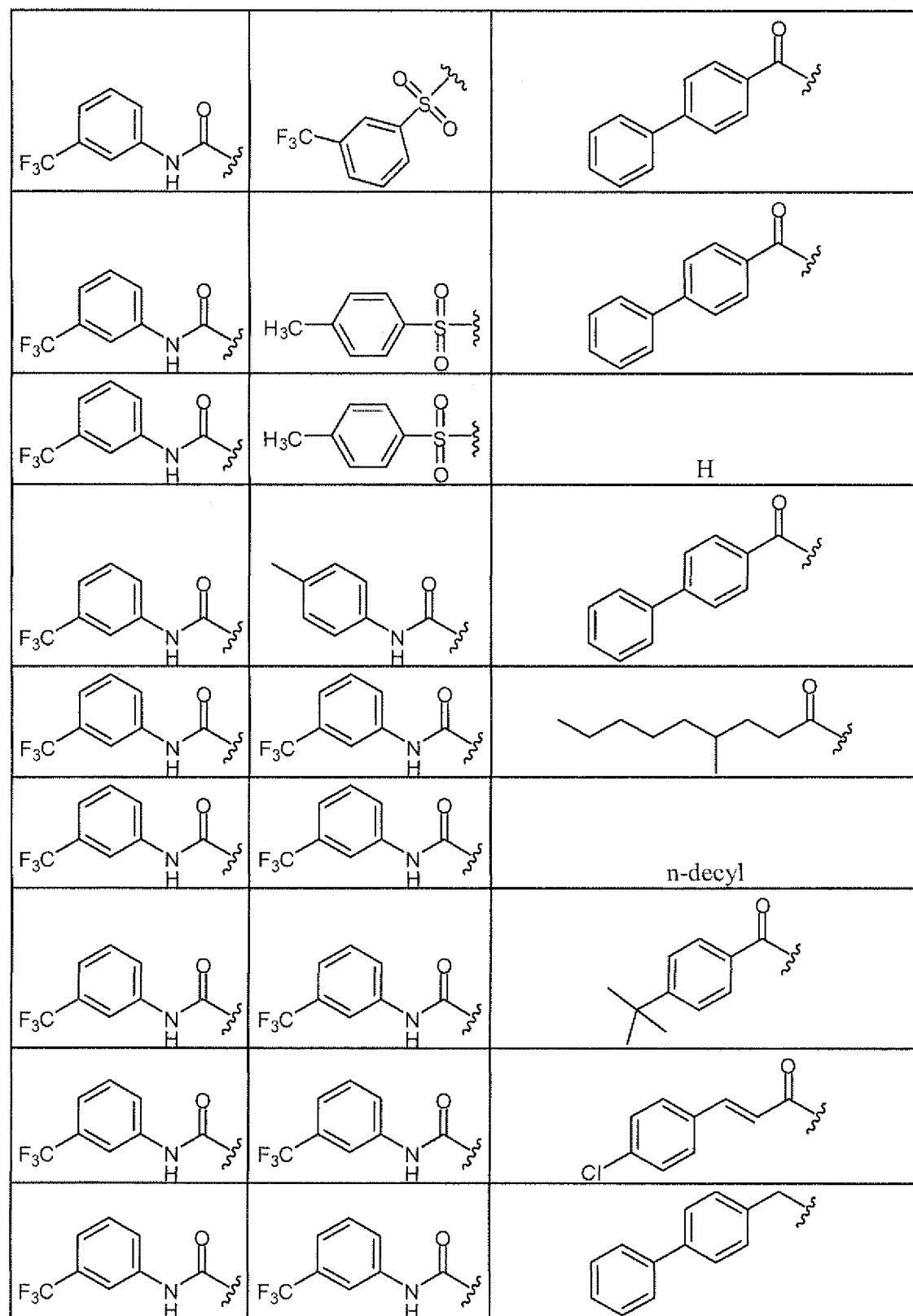
X	Y	R2	R3
CF ₃	H		
CF ₃	H		
H	CH ₃		
H	CH ₃		
H	CH ₃		

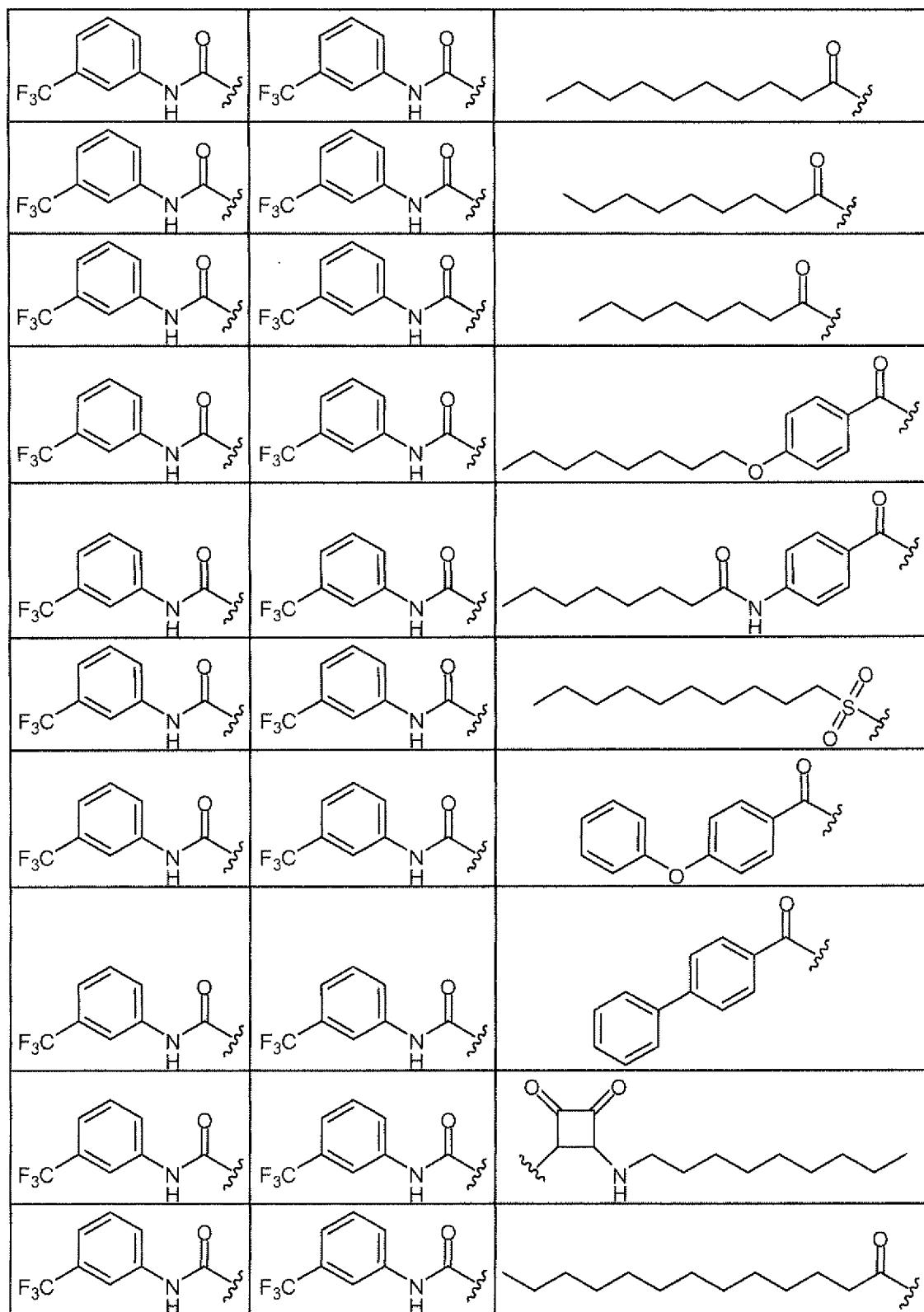
82. (New) The method of claim 70, wherein the bacteria is *Staphylococcus aureus* and the compound is selected from the group consisting of:

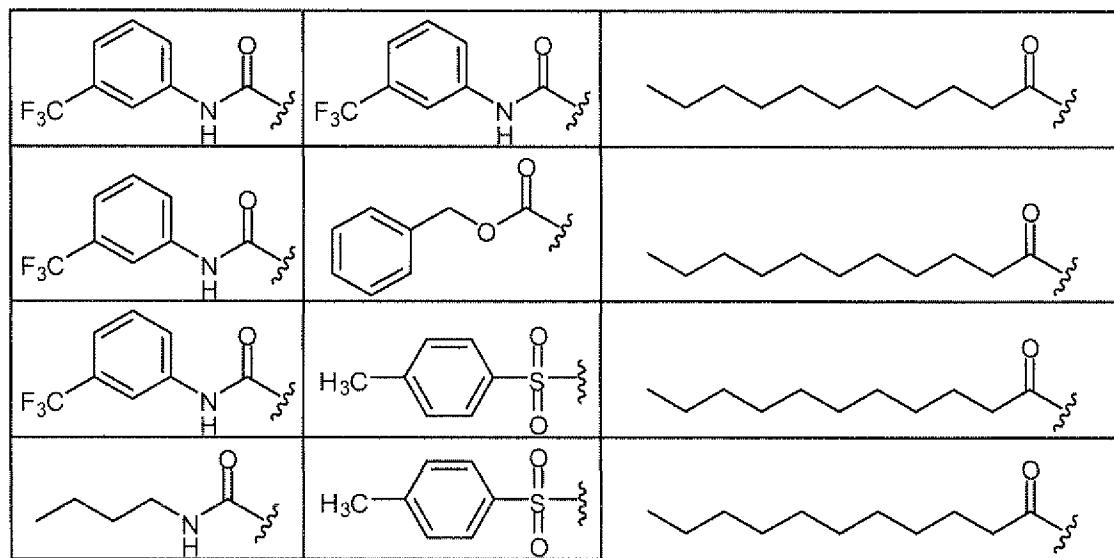


wherein

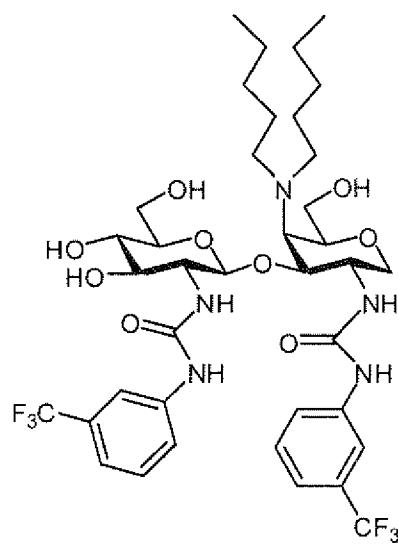
R1	R2	R3
	H	
		H
		H



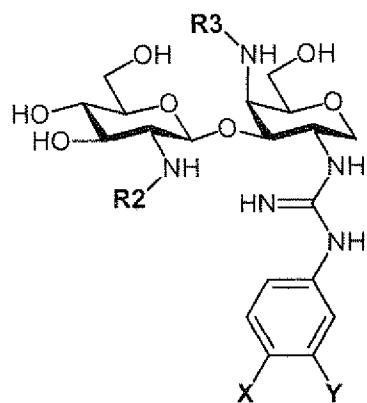




and

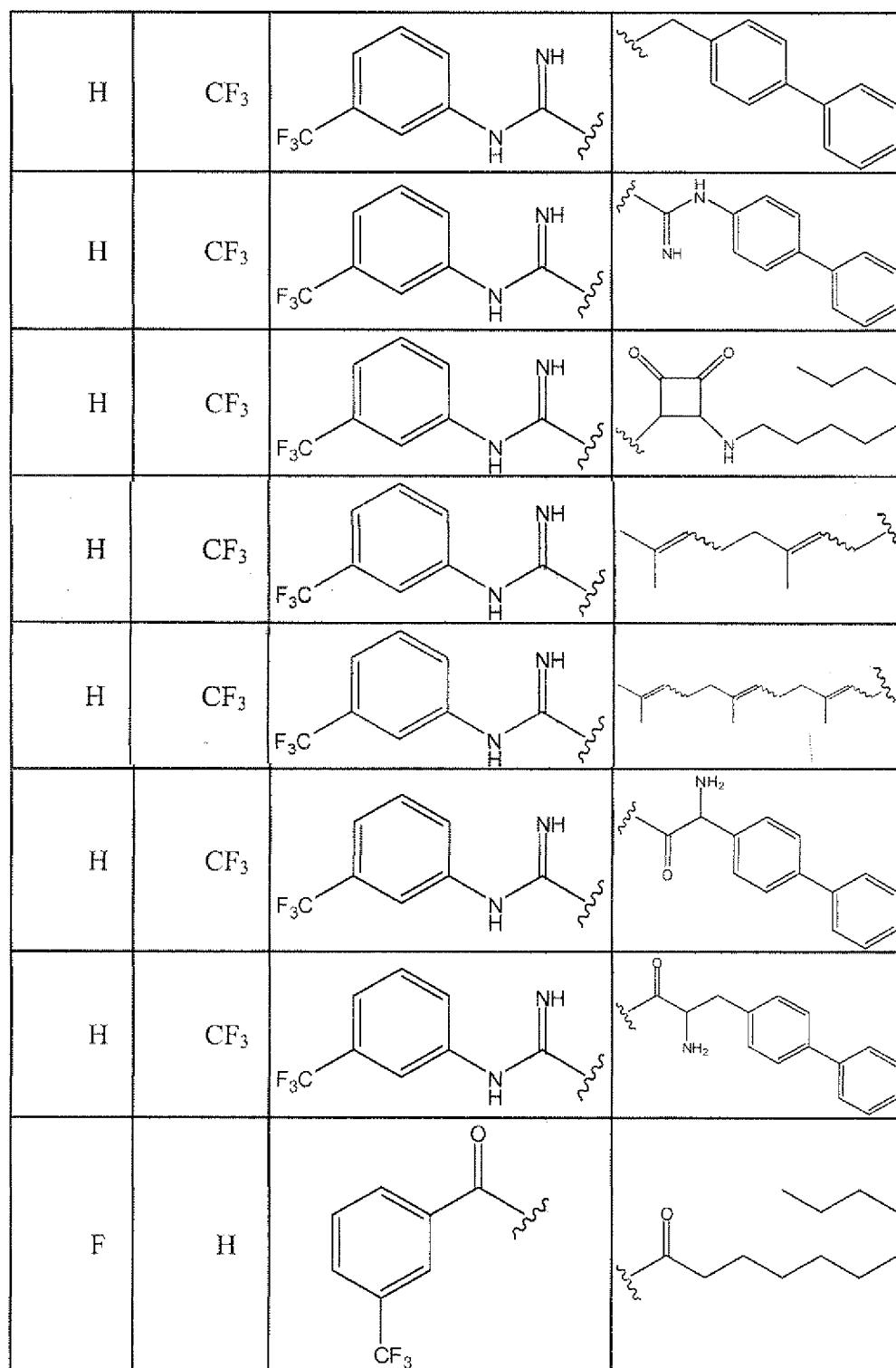


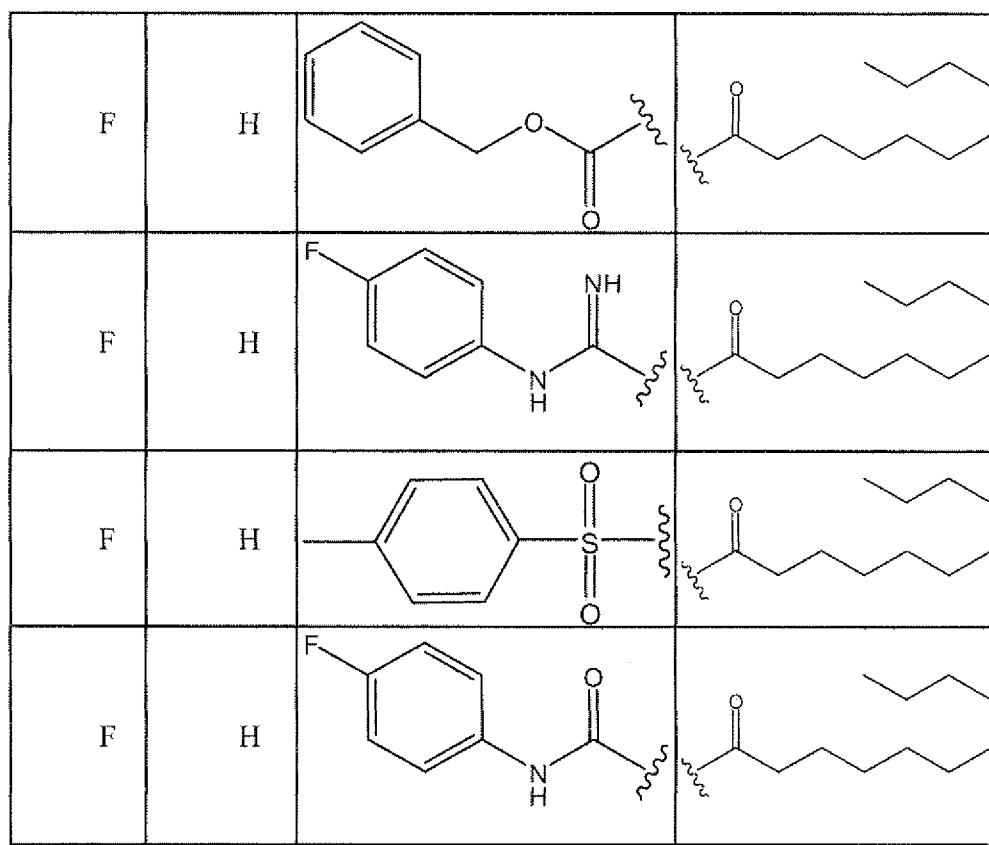
83. (New) The method of claim 70, wherein the bacteria is *E. coli* and the compound is



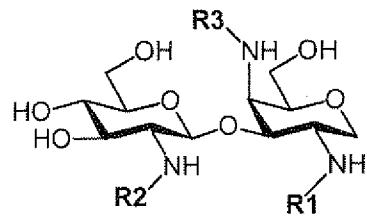
wherein:

X	Y	R2	R3
H	CF ₃		

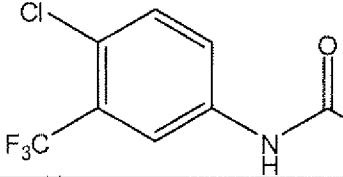
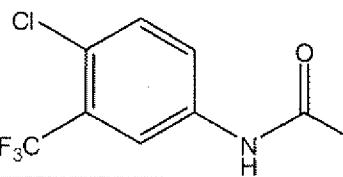
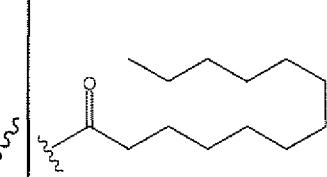


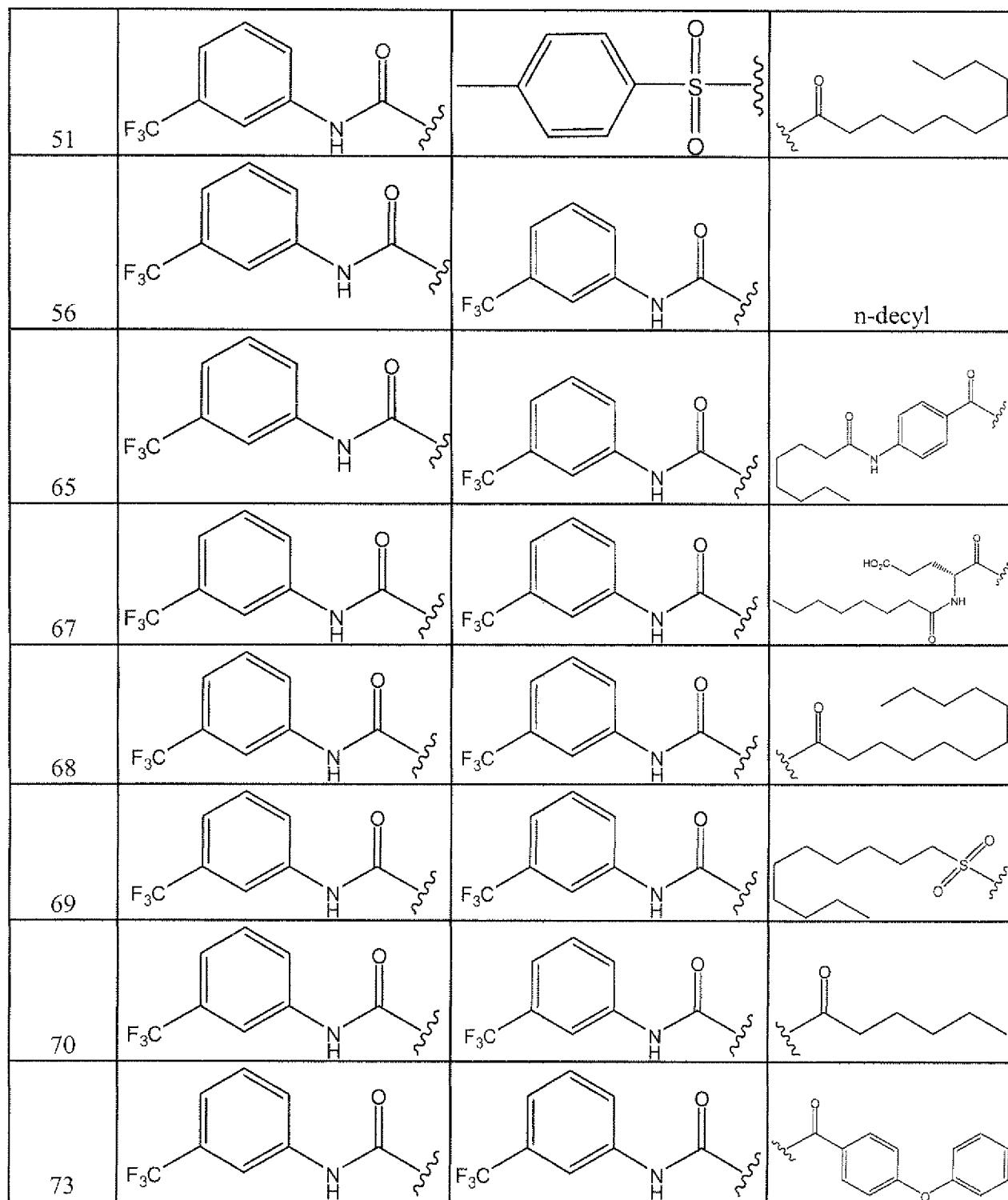


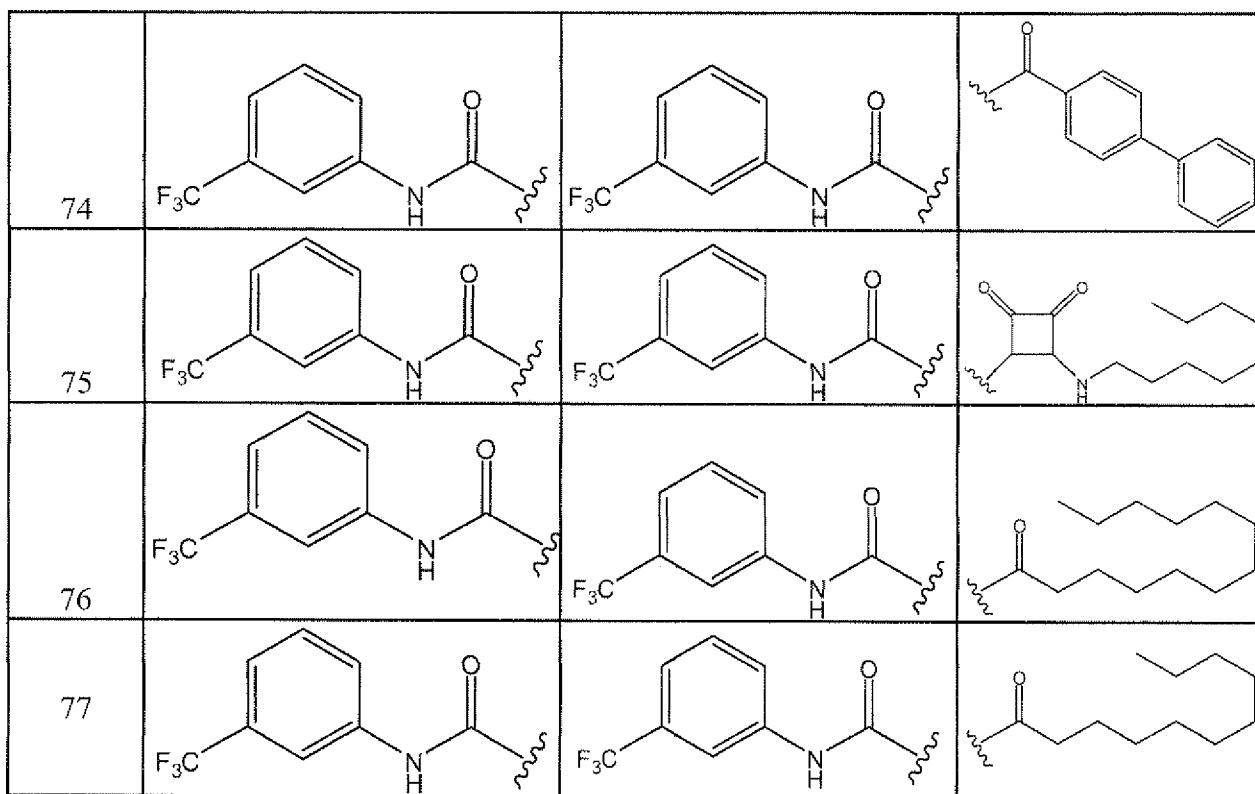
84. (New) The method of claim 70, wherein the compound is



wherein:

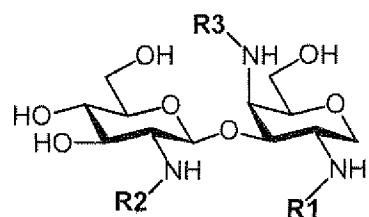
Compound	R1	R2	R3
42			



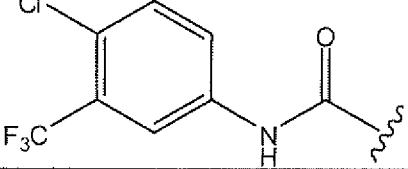
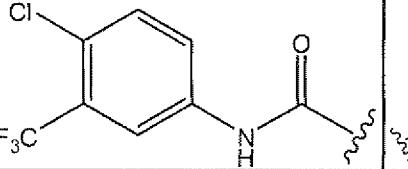
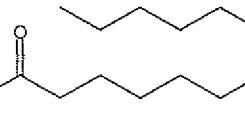
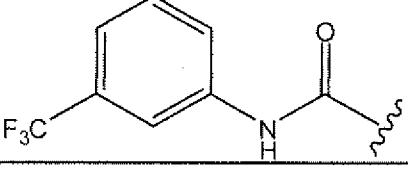
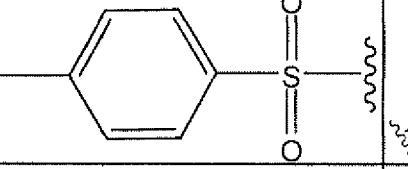
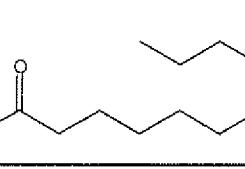
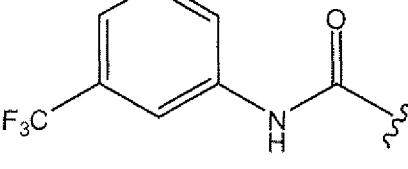
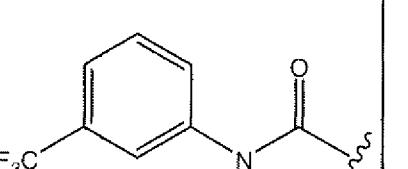
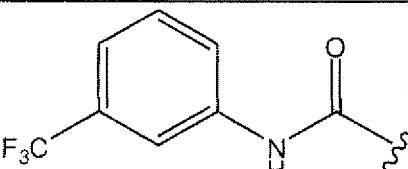
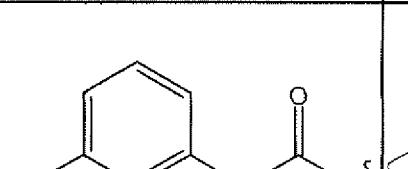
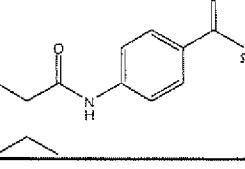
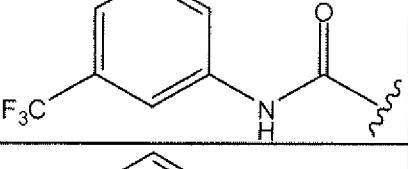
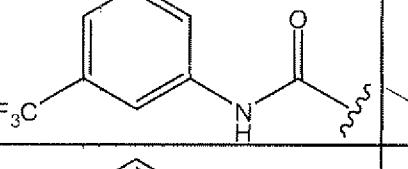
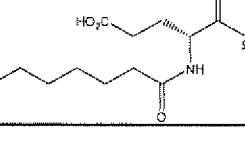
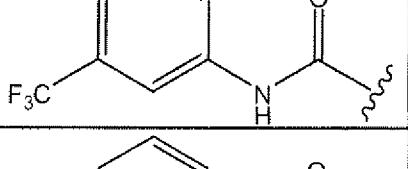
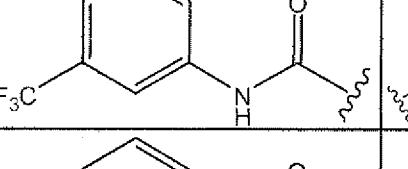
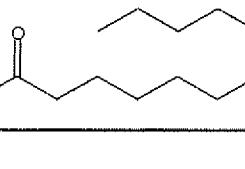
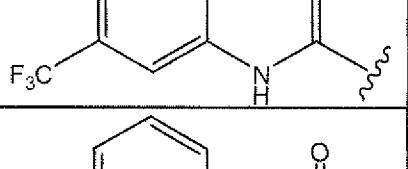
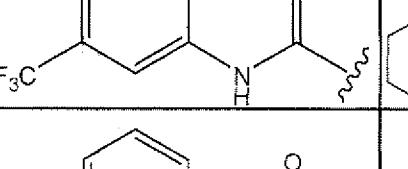
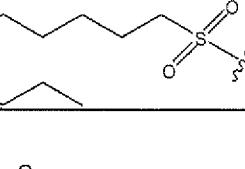
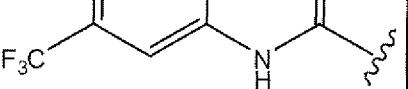
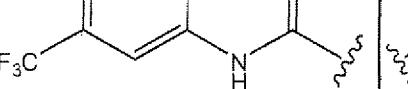
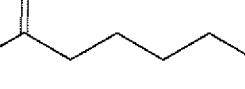


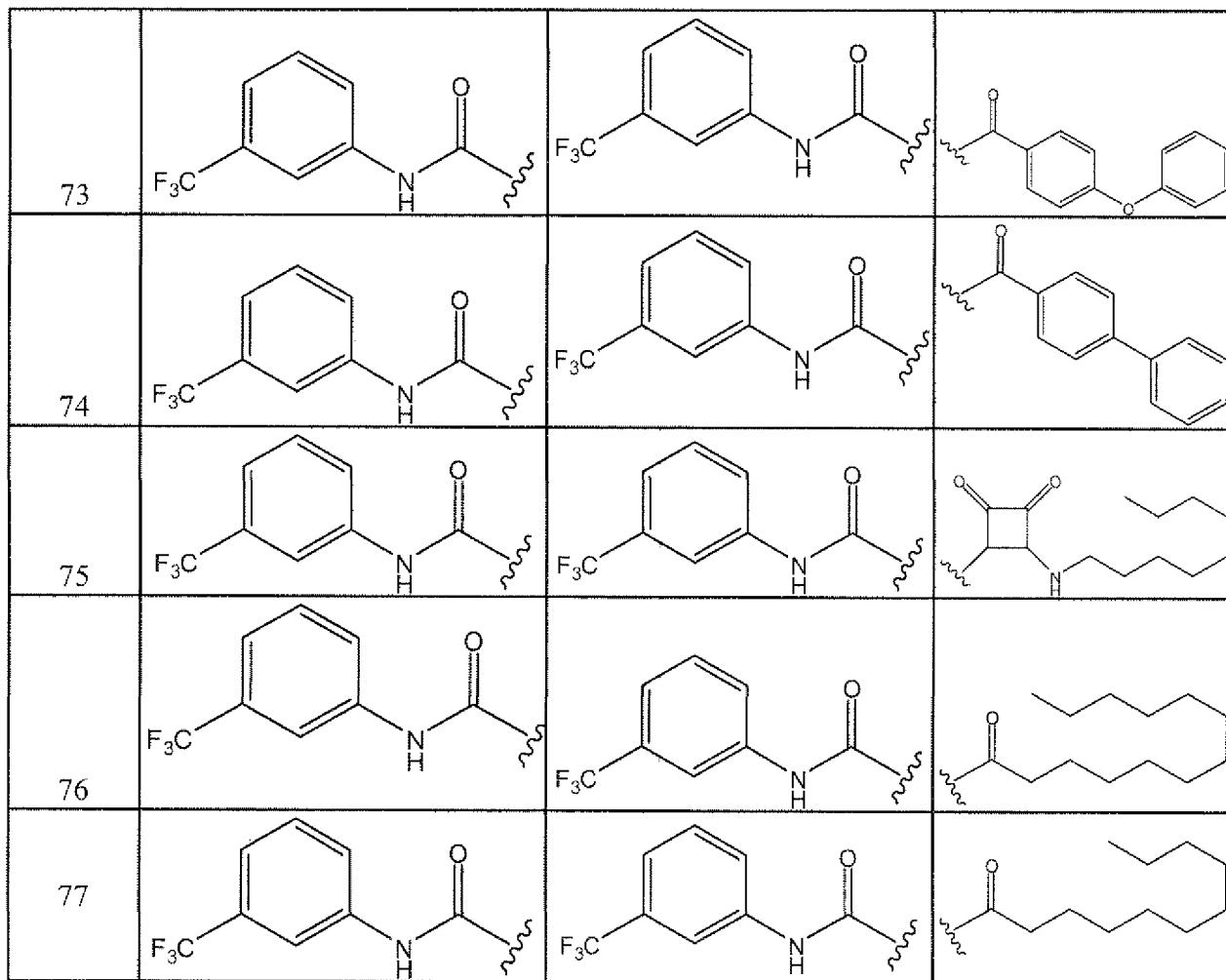
and the bacteria is *Micrococcus luteus*.

85. (New) The method of claim 70, wherein the compound is



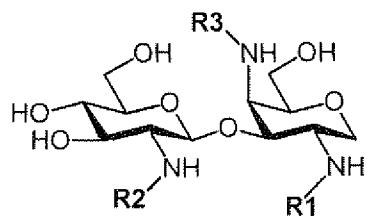
wherein:

Compound	R1	R2	R3
42			
51			
56			n-decyl
65			
67			
68			
69			
70			



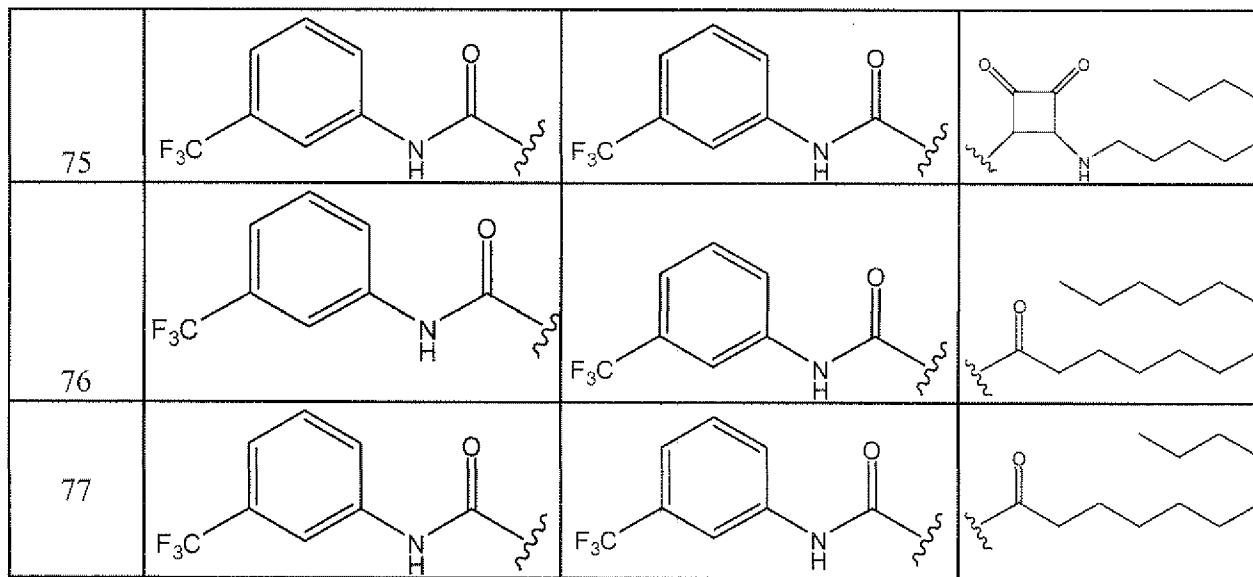
and the bacteria is *Staphylococcus aureus*.

86. (New) The method of claim 70, wherein the compound is



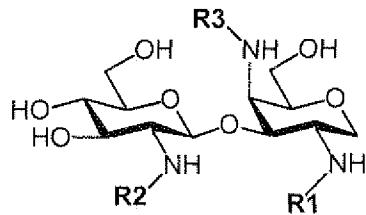
wherein:

Compound	R1	R2	R3
42			
51			
56			n-decyl
67			
68			
69			
73			
74			



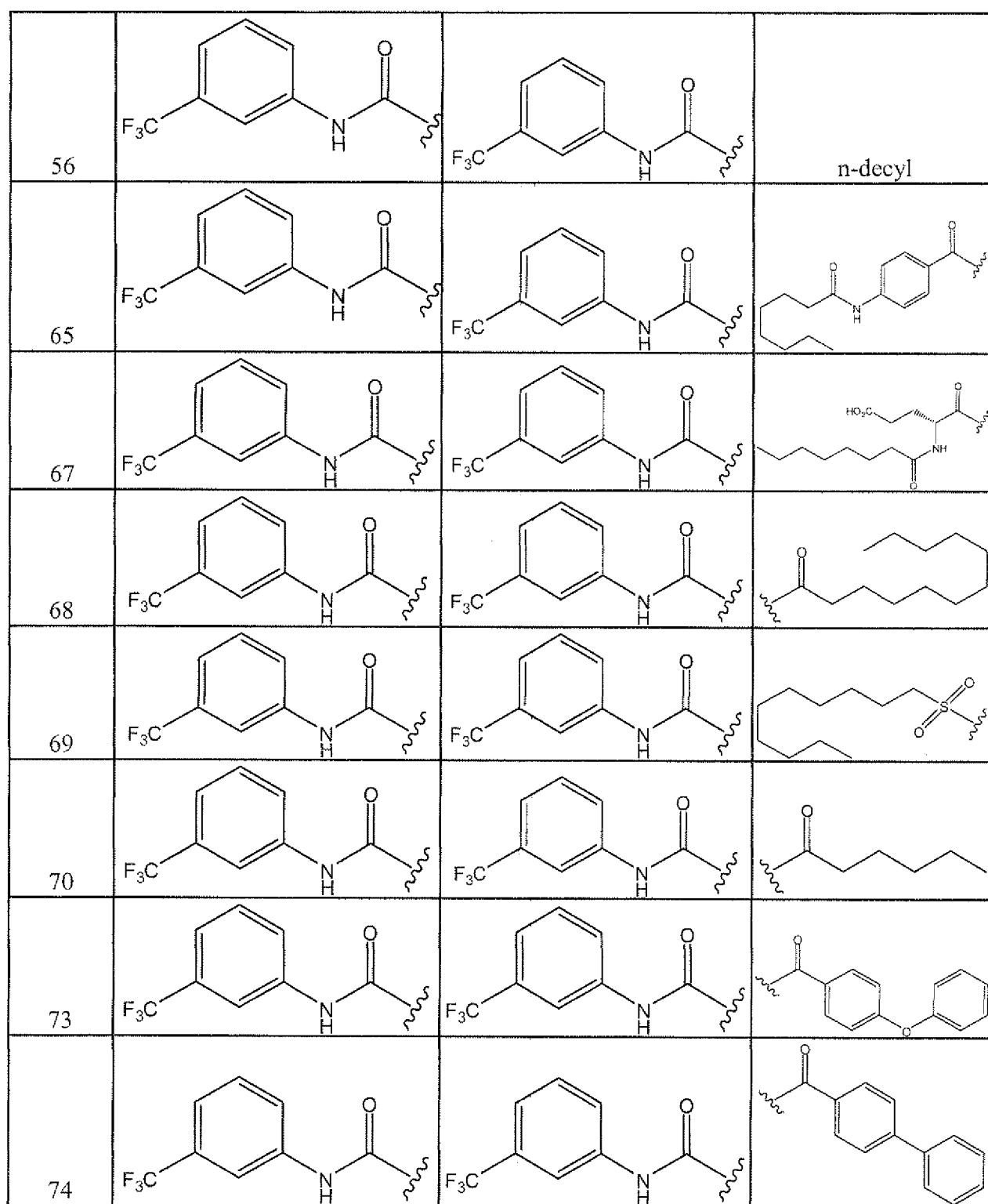
and wherein the bacteria is *Staphylococcus aureus MRS A*.

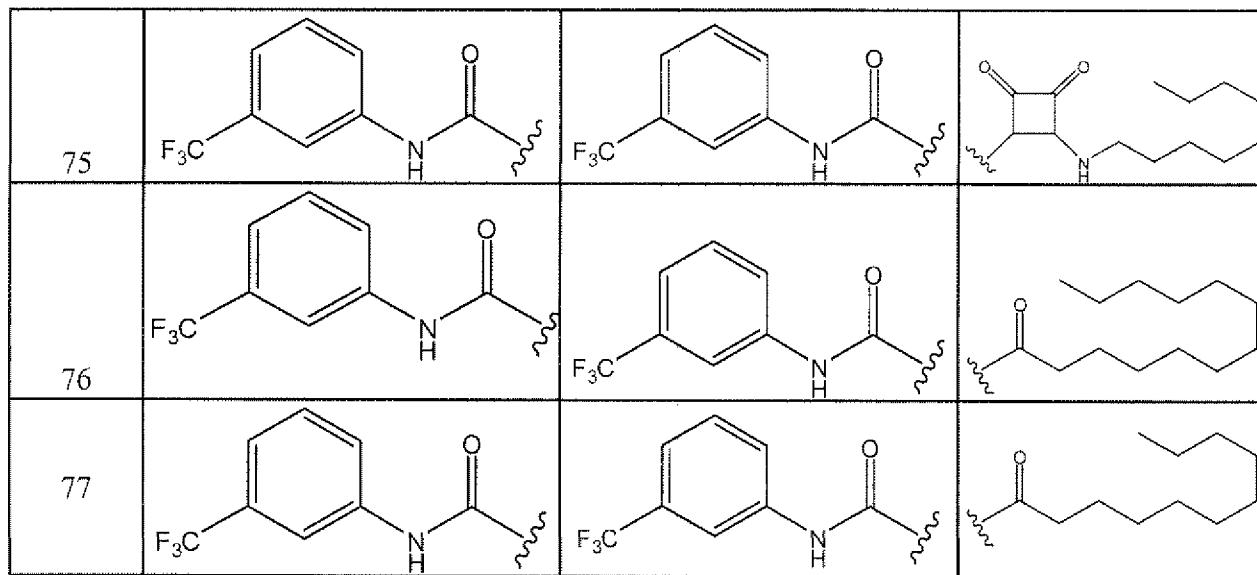
87. (New) The method of claim 70, wherein the compound is



wherein:

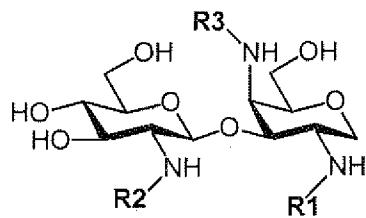
Compound	R1	R2	R3
42			
51			



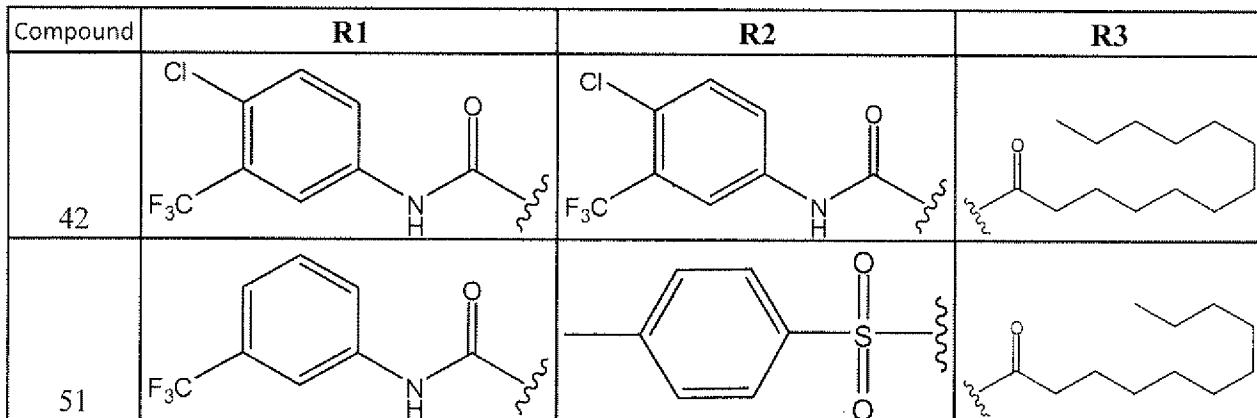


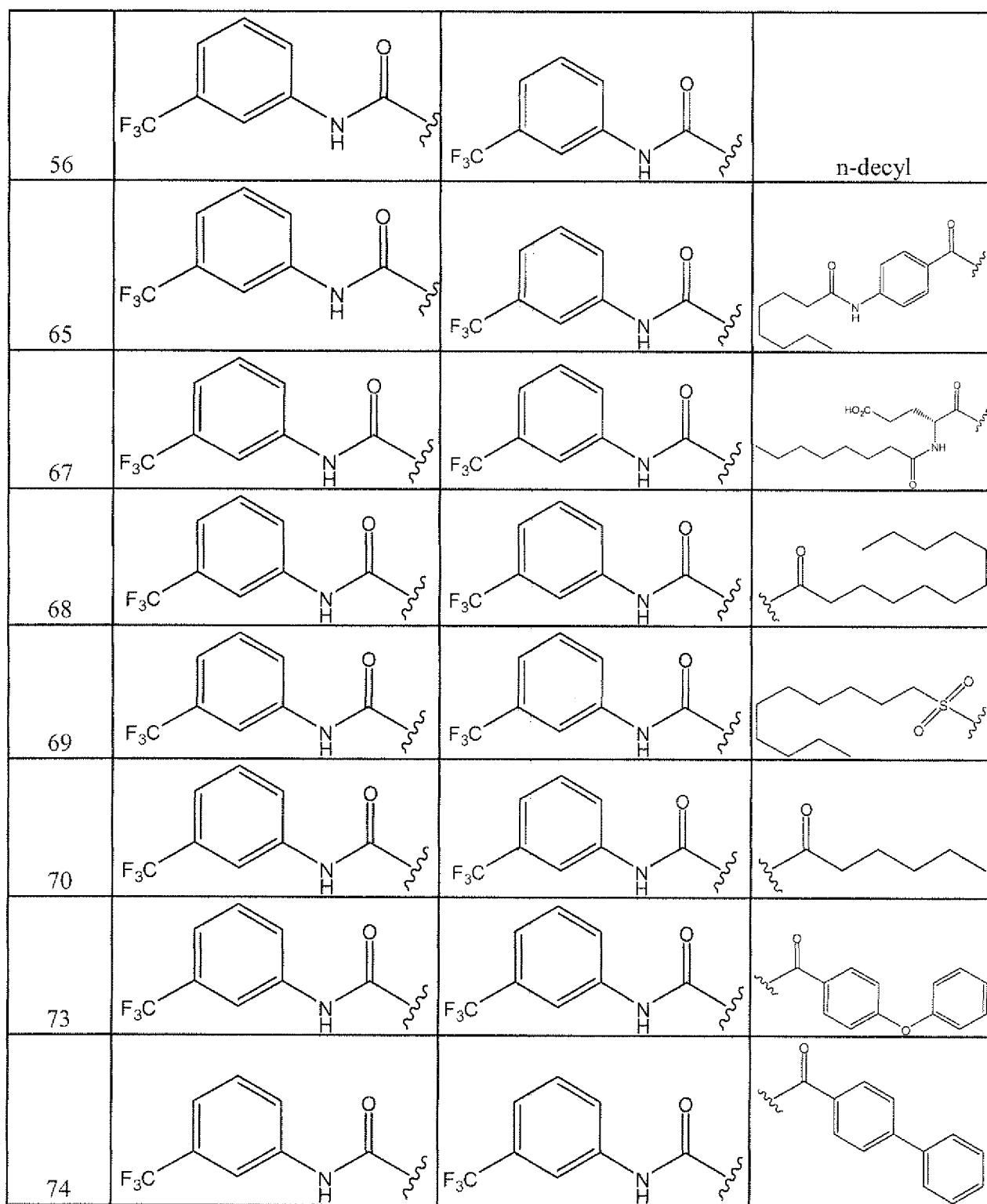
and the bacteria is *Enterococcus faecalis*.

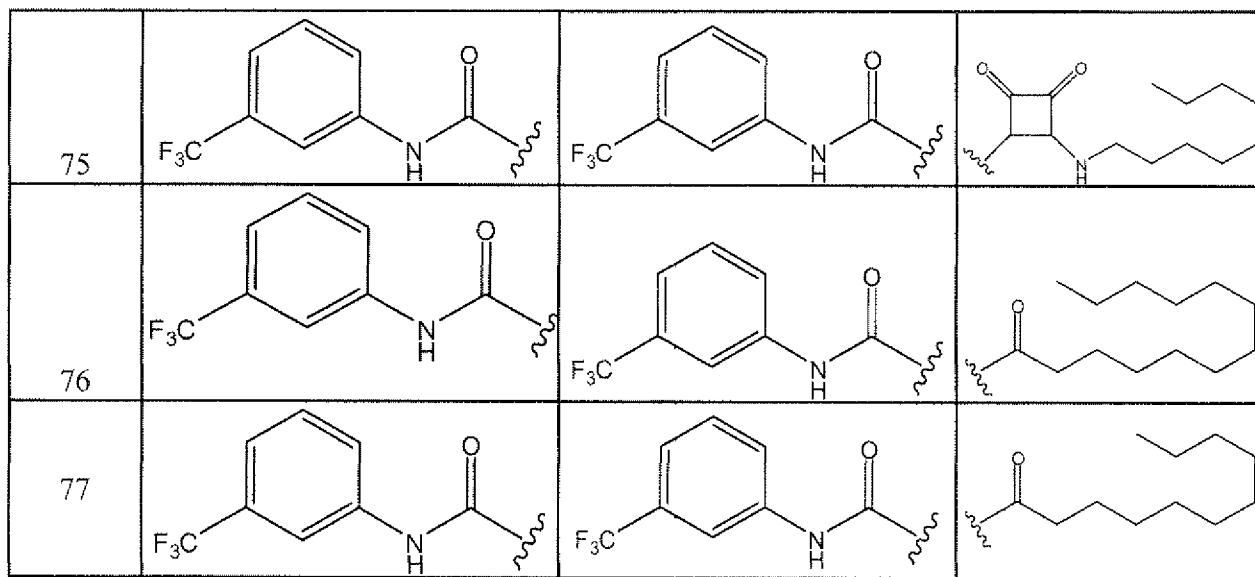
88. (New) The method of claim 70, wherein the compound is



wherein

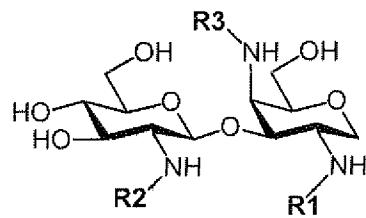




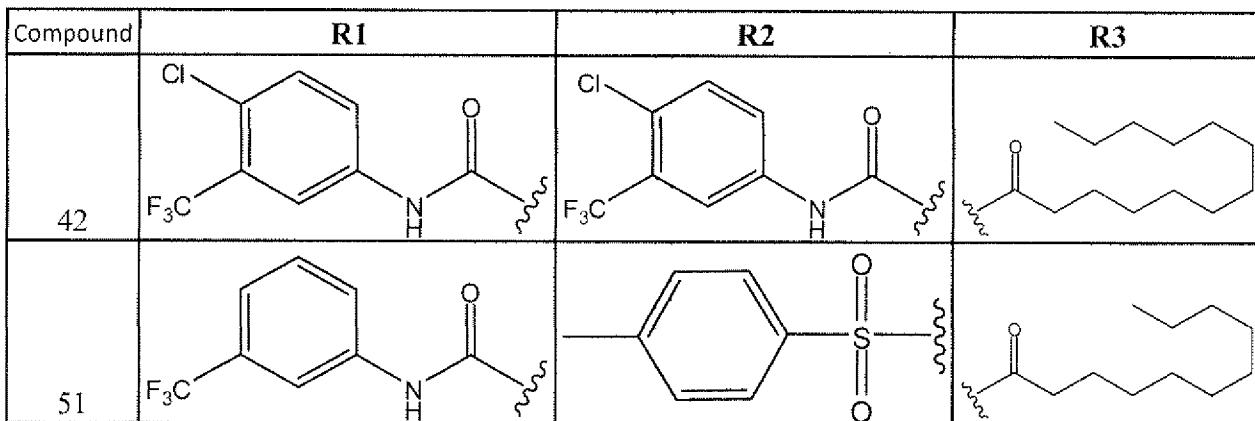


and wherein the bacteria is *Enterococcus faecalis* Vancomycin resistant.

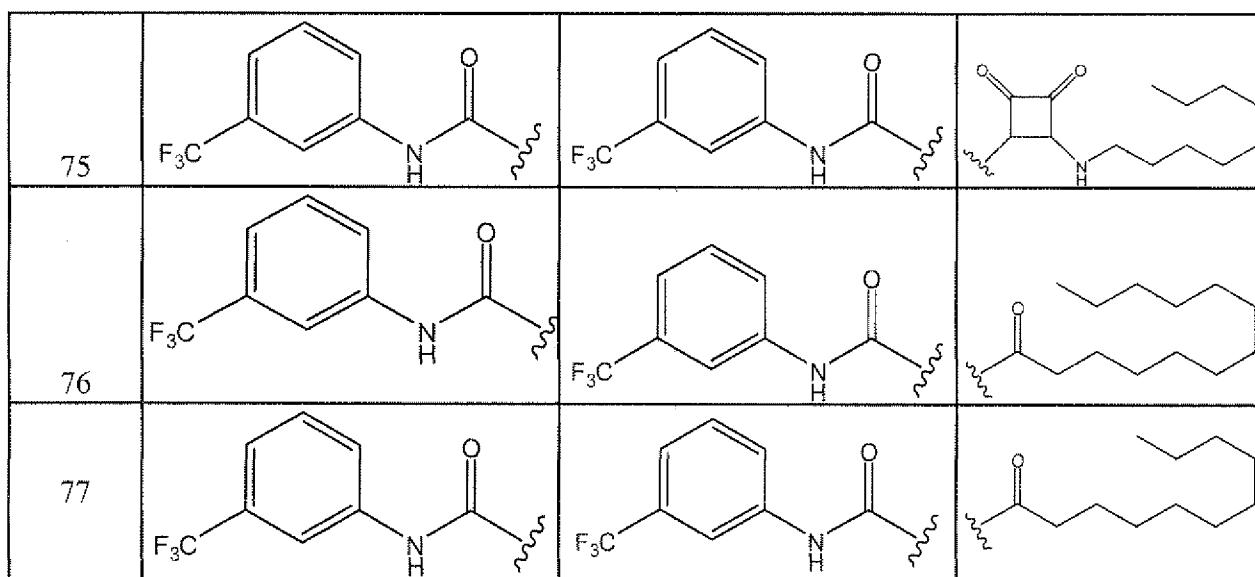
89. (New) The method of claim 70, wherein the compound is



wherein:

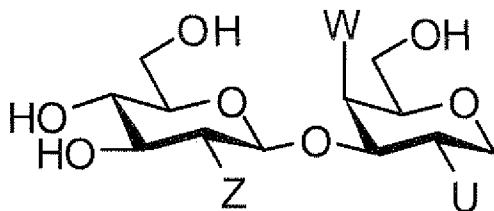


56	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	n-decyl
65	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to a propanoic acid group (-CH2-C(=O)-S).
67	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A wavy line representing a decyl chain, attached to an amide group (-NH-C(=O)-S).
68	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A wavy line representing a decyl chain, attached to a ketone group (-C(=O)-).
69	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A cyclohexane ring attached to a wavy line representing a dodecyl chain, which is attached to a thioate group (-S(=O)(=O)-S).
70	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A wavy line representing a heptyl chain, attached to a ketone group (-C(=O)-).
73	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A biphenyl ring system where the para position of one ring is attached to a phenoxy group (-O-Ph) and the para position of the other ring is attached to a carboxylic acid group (-C(=O)-COOH).
74	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A benzene ring with a trifluoromethyl group (-CF3) at the para position, attached to an amide group (-NH-C(=O)-S).	A biphenyl ring system where the para position of one ring is attached to a phenyl group (-Ph) and the para position of the other ring is attached to a carboxylic acid group (-C(=O)-COOH).



and the bacteria is *Streptococcus pyogenes*.

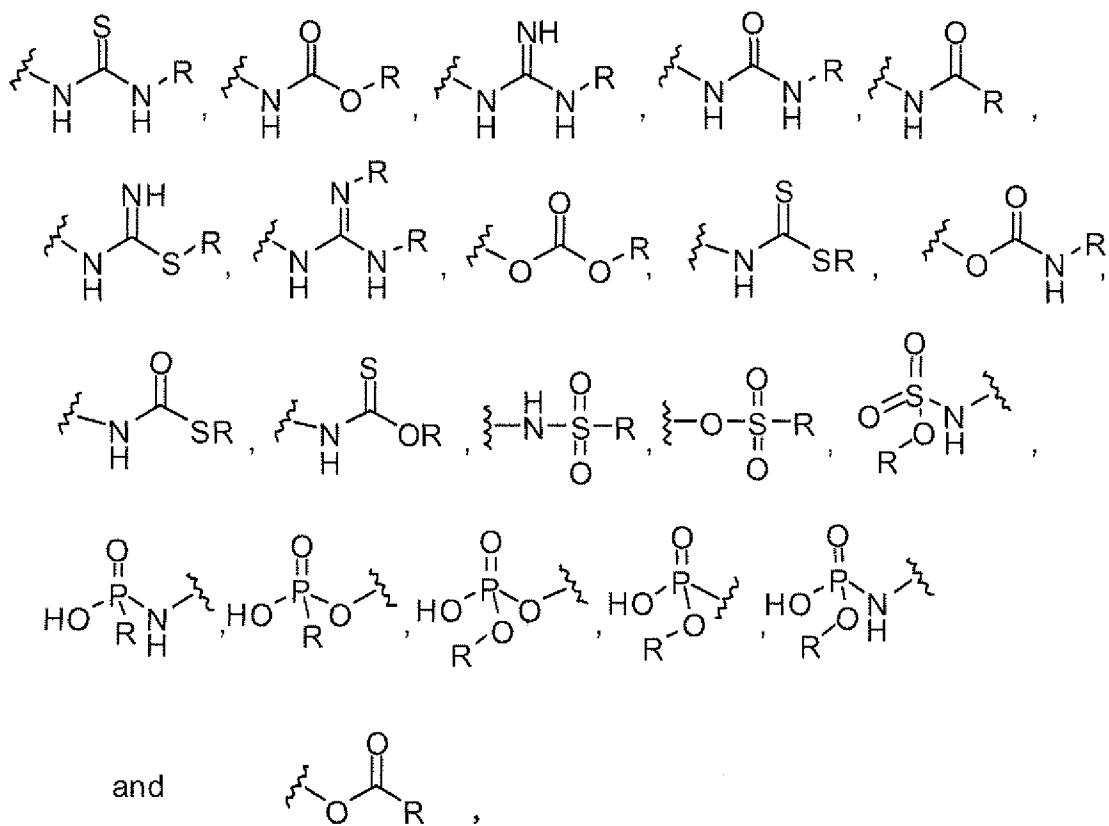
90. (New) A method of inhibiting a bacterial infection in a mammal comprising administering to said mammal an effective amount of a compound of General Formula (I),



General Formula (I)

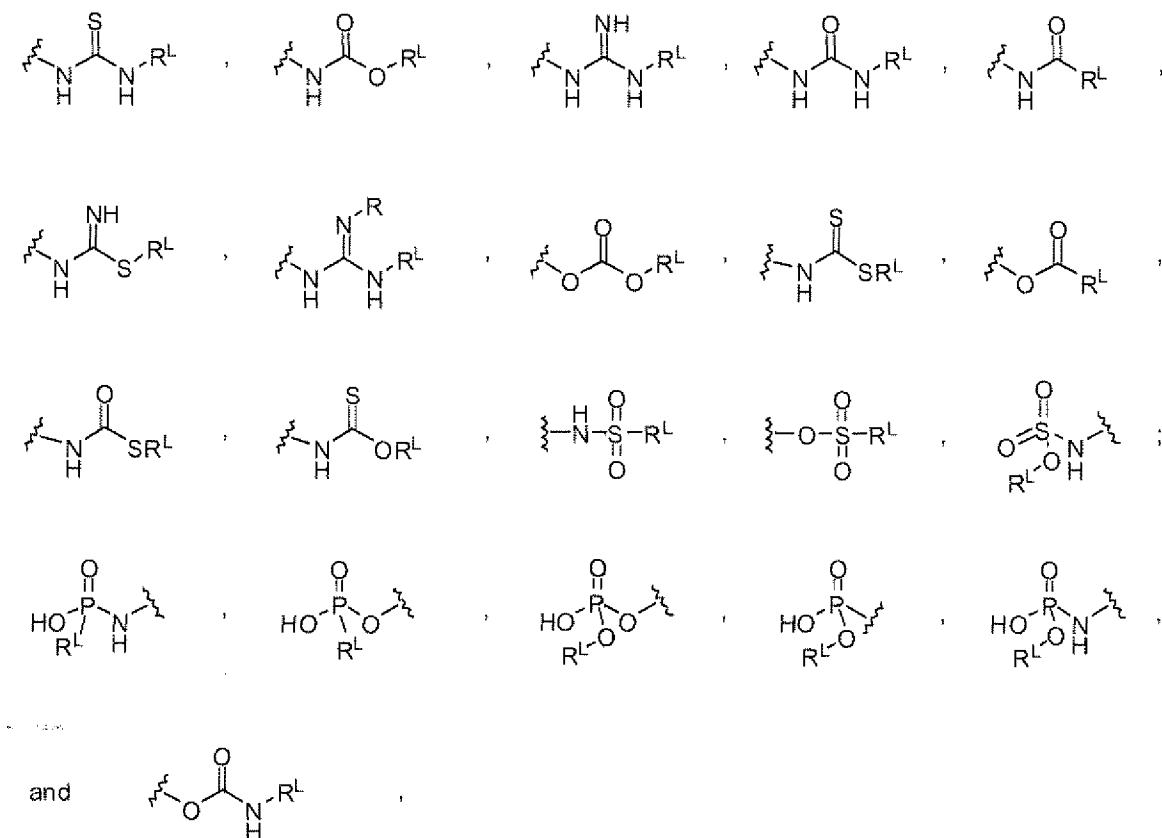
wherein

U and Z are independently selected from the group consisting of: -OR, -NHR, and -NR(R),



wherein R may be the same or different, R is a moiety of not more than 20 carbon atoms independently selected from the group consisting of: alkyl, alkenyl, alkynyl, heteroalkyl, aryl, heteroaryl, arylalkyl and heteroarylalkyl;

W is independently selected from the group consisting of - OR^L , - NHR^L , - NR^LR ,



wherein R^L is a substituted or unsubstituted, linear or branched moiety of between 3 and 55 carbon atoms selected from the group consisting of: alkyl, heteroalkyl, arylalkyl, and alkylaryl chain.

91. (New) The method of claim 70, wherein the bacterium is a resistant or susceptible strain of a *Micrococcus*, *Streptococcus*, *Enterococcus* or *Staphylococcus*.